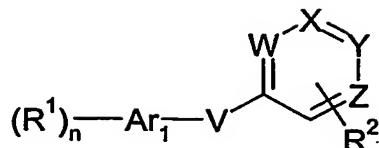


CLAIMS

1. A compound of formula (I):



(I)

wherein

V represents NR⁵, O, S, SO or S(O)₂;

W and X each independently represent CH or N;

Y represents N, CH or C-Ar₂, with the proviso that at least one, but no more than two, of W,

10 X and Y are N;

Z represents CH or C-Ar₂, with the proviso that when Y is N or CH then Z is C-Ar₂, and with the further proviso that when Y is C-Ar₂ then Z is CH;

Ar₁ represents a fused 9 or 10 membered heterobicyclic ring system containing one, two, three or four heteroatoms selected from nitrogen, oxygen and sulfur, wherein at least one of the 15 rings in said ring system is aromatic;

Ar₂ represents an aromatic ring selected from phenyl, pyridyl, pyridazinyl, pyrimidinyl and pyrazinyl; which aromatic ring is optionally fused to a phenyl ring, a five-membered heteroaromatic ring containing 1, 2, 3 or 4 heteroatoms selected from O, N and S at most 1 heteroatom being O or S, or a six-membered heteroaromatic ring containing 1, 2 or 3 N atoms; which aromatic ring is unsubstituted or substituted by one, two or three groups selected from halogen, hydroxy, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenylC₁₋₂alkoxy, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy, haloC₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₃₋₇cycloalkyl, C₃₋₇cycloalkoxy, C₃₋₅cycloalkylC₁₋₄alkyl, cyano, nitro, SR⁶, SOR⁶, SO₂R⁶, COR⁶, NR³COR⁶, CONR³R⁴, NR³SO₂R⁶, SO₂NR³R⁴, -(CH₂)_mcarboxy, esterified

20 -(CH₂)_mcarboxy, -(CH₂)_mNR³R⁴, phenyl, naphthyl, a five-membered heteroaromatic ring containing 1, 2, 3 or 4 heteroatoms selected from O, N and S at most 1 heteroatom being O or S and a six-membered heteroaromatic ring containing 1, 2 or 3 N atoms; where two C₁₋₆alkoxy groups are on adjacent atoms they may, together with the atoms to which they are attached, form a 5- or 6-membered partially saturated ring;

30 R¹ represents halogen, hydroxy, oxo, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy, haloC₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₃₋₇cycloalkyl, C₃₋₇cycloalkoxy.

γ cycloalkoxy, C_{3-5} cycloalkyl C_{1-4} alkyl, cyano, nitro, SR^6 , SOR^6 , SO_2R^6 , COR^6 , NR^3COR^6 , $CONR^3R^4$, $NR^3SO_2R^6$, $SO_2NR^3R^4$, $-(CH_2)_m$ carboxy, esterified $-(CH_2)_m$ carboxy or $-(CH_2)_mNR^3R^4$;

R^2 represents hydrogen, halogen, hydroxy, C_{1-6} alkyl, halo C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{1-6} alkoxy,

5 halo C_{1-6} alkoxy, unsubstituted phenyl or phenyl substituted with one or two groups selected from halogen, C_{1-6} alkyl, halo C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{1-6} alkoxy or halo C_{1-6} alkoxy;

R^3 and R^4 are each independently hydrogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{3-7} cycloalkyl or fluoro C_{1-6} alkyl;

or R^3 and R^4 and the nitrogen atom to which they are attached together form a heteroaliphatic

10 ring of 4 to 7 ring atoms, optionally substituted by one or two groups selected from hydroxy or C_{1-4} alkoxy, which ring may optionally contain as one of the said ring atoms an oxygen or a sulfur atom, $S(O)$, $S(O)_2$, or NR^5 ;

R^5 represents hydrogen, C_{1-4} alkyl, hydroxy C_{1-4} alkyl or C_{1-4} alkoxy C_{1-4} alkyl;

R^6 represents hydrogen, C_{1-6} alkyl, fluoro C_{1-6} alkyl, C_{3-7} cycloalkyl, unsubstituted phenyl, or

15 phenyl substituted with one or two groups selected from halogen, C_{1-6} alkyl, halo C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{1-6} alkoxy or halo C_{1-6} alkoxy;

m is either zero or an integer from 1 to 4;

n is either zero or an integer from 1 to 3;

or a pharmaceutically acceptable salt, N -oxide or a prodrug thereof.

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2. A compound according to claim 1 in which R^1 is halogen, C_{1-4} alkyl or fluoro C_{1-4} alkyl.

3. A compound according to claim 1 or 2 in which n is one or two.

25 4. A compound according to claim 1, 2 or 3 in which R^2 is hydrogen, halogen, C_{1-4} alkyl, C_{1-4} alkoxy or phenyl substituted by C_{1-4} alkyl or fluoro C_{1-4} alkyl.

5. A compound according to any preceding claim in which $=W-X=Y-$ represents $=N-CH=CH-$, $=N-N=CH-$, $=N-CH=N-$ or $=N-N=C(Ar_2)-$.

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6. A compound according to any preceding claim in which Ar_1 represents a heterobicyclic ring system selected from isoquinoline, indazole, triazolopyridine, cinnoline, benzothiazole, imidazopyridine, quinoline, tetrahydroisoquinoline or dihydroisoquinoline.

35 7. A compound according to any preceding claim in which Ar_2 is phenyl or pyridyl which are optionally fused to a phenyl, imidazolyl or thienyl ring, and are unsubstituted or

substituted by one to three groups independently selected from halogen, cyano, C₁₋₄alkyl, fluoroC₁₋₄alkyl, C₁₋₄alkoxy, fluoroC₁₋₄alkoxy, phenylC₁₋₂alkoxy, piperidine optionally substituted by oxygen, COR⁶ where R⁶ is hydrogen or C₁₋₄alkyl, pyrazole, C₁₋₄alkylcarbonyl, carboxy, C₁₋₆alkylsulphonyl, nitro, phenyl, C₁₋₄alkylthio, hydroxy and -O-CH₂-O-.

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8. A pharmaceutical composition comprising a compound of formula (I) according to any preceding claim, or a pharmaceutically acceptable salt or N-oxide thereof, and a pharmaceutically acceptable excipient.

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9. A compound of formula (I) according to any one of claims 1 to 7, or a pharmaceutically acceptable salt or N-oxide thereof, for use in a method of treatment of the human or animal body by therapy.

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10. Use of a compound of formula (I) according to any one of claims 1 to 7, or a pharmaceutically acceptable salt or N-oxide thereof, for use in the manufacture of a medicament for the treatment or prevention of a disease or condition in which pain and/or inflammation predominates.

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11. A method for the treatment or prevention of a disease or condition in which pain and/or inflammation predominates, which method comprises administration to a patient in need thereof of an effective amount of a compound of formula (I) according to claim 1, or a pharmaceutically acceptable salt or N-oxide thereof.